

CLAIMS

1. A method for preparing a drug eluting medical device comprising the application to said device of a polymer having active functional groups capable of chemically
5 binding biological molecules, characterised in that said application takes place in a single step by means of cold plasma methods.
2. A method according to claim 1, in which said polymers are chosen from among polymers having amine groups,
10 carboxyl groups and sulphhydryl groups.
3. A method according to claim 2 in which the precursors of said polymers having amine groups are chosen from among allylamine, heptylamine, aliphatic amines and aromatic amines.
- 15 4. A method according to claim 2 in which the precursors of said polymers having carboxylic groups are chosen from between acrylic acid and methacrylic acid.
5. A method according to claim 2, in which the precursors of said polymers having sulphhydryl groups are chosen
20 from among volatile mercaptans.
6. A method according to claim 1, in which said cold plasma methods comprise cold plasma produced under vacuum using discontinuous or continuous technology.
7. A method according to claim 6, in which said cold
25 plasma under vacuum is generated at a pressure which may

vary between 0.01 and 10 mbar, at a power of between 1 and 500 W and for a period of time of not more than 30 minutes.

8. A method according to claim 1, in which said cold
5 plasma methods consist in cold plasma produced at atmospheric pressure.

9. A method according to claim 1 in which the precursor of said polymer is in the form of a gas.

10. A method according to claim 1, in which the precursor
10 of said polymer is in the form of a vapour.

11. A method according to claim 1, in which said polymer is applied in the form of film with a thickness of between 0.01 and 10 microns.

12. A method according to claim 1, also comprising before
15 the application of said polymer having functional groups a step of applying at least one layer of a drug incorporated where appropriate in a polymer capable of eluting said drug.

13. A method according to claim 12, in which said drug is
20 chosen from the group consisting in anti-inflammatory, anti-proliferative and anti-migratory drugs and immunosuppressive agents.

14. A method according to claim 13, in which said drug is
4-[(4-methyl-1-piperazinyl)methyl]-N-[4-methyl-3-[[4-(3-
25 pyridinyl)-2-pyrimidinyl]amino]-phenyl]benzamide

methanesulphonate.

15. A method according to claim 12, in which the drug
eluting polymer is chosen from among hydrophobic
hydrocarbons, polyamides, polyacrylates and
5 polymethacrylates.

16. A method according to claim 15 in which said
hydrophobic hydrocarbons are chosen from among
polystyrene, polyethylene, polybutadiene and
polyisoprene.

10 17. A method according to claim 15, in which said polymer
is chosen from among polyhydroxybutylmethacrylate,
polyhydroxyethylmethacrylate, where appropriate in
combination with polybutadiene.

18. A method according to claim 12 in which said drug
15 which may be incorporated in a drug eluting polymer is
applied by means of immersion in a suitable solution or
deposited by spraying.

19. A method according to claim 18 in which said drug
eluting polymer is deposited in the form of film with a
20 thickness of between 0.5 and 20 microns.

20. A method according to claim 12, in which when said
drug is an anti-inflammatory, it is present in quantities
of between 0.001 mg and 10 mg per device.

21. A method according to claim 12, in which when said
25 drug is an anti-proliferative, it is present in

quantities of between 0.0001 and 10 mg per device.

22. A method according to claim 12, in which when said drug has an anti-migratory action, it is present in quantities of between 0.0001 mg and 10 mg per device.

5 23. A method according to claim 12, in which when the drug is an immunosuppressant, it is present in quantities of between 0.0001 mg and 10 mg per device.

24. A method according to claims 1 in which when said drug is 4-[(4-methyl-1-piperazinyl)methyl]-N-[4-methyl-3-
10 [[4-(3-pyridinyl)-2-pyrimidinyl]amino]-phenyl]benzamide methanesulphonate, it is present in quantities of between 0.001 mg and 10 mg per device.

25. A method according to claim 1, also comprising a step of depositing biological molecules on the surface of said
15 polymer having stable reactive functional groups.

26. A method according to claim 25, in which said biological molecules are chosen from among anti-thrombotic substances and hyaluronic acid.

27. A method according to claim 26, in which said
20 biological molecules are heparin.

28. A method according to claim 26, in which said biological molecules are deposited by immersing the medical device in an aqueous solution containing said biological molecules in a concentration of 0.01% to 1% by
25 weight.

29. A method according to claim 1, also comprising a preliminary step of cleaning/washing said medical device.

30. A method according to claim 29, in which said preliminary cleaning/washing step is followed by a step
5 of pretreatment of said medical device to promote adhesion of the drug incorporated where appropriate in an eluting polymer to this device.

31. A method according to claim 1, also comprising the application of further biodegradable polymer layers over
10 said biological molecule layer.

32. A method according to claim 1, comprising in succession the application of at least one first layer of 4-[(4-methyl-1-piperazinyl)methyl]-N-[4-methyl-3-[[4-(3-pyridinyl)-2-pyrimidinyl]amino]-phenyl]benzamide
15 methanesulphonate included where appropriate in a polymer to the surface of said medical device, the application by cold plasma of at least one second layer of polymer of allylamine, the bonding of heparin to said at least one second layer and application of at least one third layer
20 of biodegradable polymer onto said heparin.

33. A drug eluting medical device obtainable by means of the method according to any one of the preceding claims.

34. A medical device according to claim 33, comprising a device structure, at least one first layer covering the
25 surface of said structure comprising a drug, at least one

second layer covering said at least one first layer comprising a polymer having stable reactive functional groups and a biological molecule layer bonded to said at least one second layer by means of chemical bonding with
5 said functional groups, in which said at least one second layer of polymer is deposited on said at least one first layer by means of a cold plasma method.

35. A medical device according to claim 34, in which said drug is a drug as described in any one of claims 13 to
10 32.

36. A medical device according to claim 34, in which said drug eluting polymer is a polymer as described in claim 16.

37. A medical device according to claim 34, in which said
15 polymer having stable reactive functional groups is one of the polymers described in claim 2.

38. A medical device according to claim 34, in which said biological molecule is any one of the molecules in claim 26.

20 39. A medical device according to claim 34, said device being chosen from among vascular devices, prostheses, probes, catheters, dental implants or similar.

40. A medical device according to claim 39, said device being a vascular stent.

25 41. The use of polymers having reactive functional groups

chosen from among the polymers described in claim 2, for covering medical devices, preferably vascular stents, by means of cold plasma methods of deposition.